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STRUCTURE UPLOADED

=> s 19 sss full

FULL SEARCH INITIATED 17:48:08 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 378141 TO ITERATE

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 12 ANSWERS

 60.1% PROCESSED
 227393 ITERATIONS
 54 ANSWERS

 82.3% PROCESSED
 311374 ITERATIONS
 59 ANSWERS

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 69 ANSWERS

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 70 ANSWERS

L11 70 SEA SSS FUL L9

=> d 19 sim

L9 HAS NO ANSWERS L9 STI

SEARCH TIME: 00.01.22

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT * Structure attributes must be viewed using STN Express query preparation.

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14531335 UP<19981130 (UP<981130) 17 L11 AND UP<19981130

=> s pyrid?/cns and l11

1098272 PYRID?/CNS L13 38 PYRID?/CNS AND L11

=> s 113 not 112

L12

L14 38 L13 NOT L12

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L12 ANSWER 1 OF 17 REGISTRY COPYRIGHT 2000 ACS
RN 188811-83-8 REGISTRY
CN .beta.-Alanine, N-[3-[(aminoiminomethyl)amino]benzoyl]glycyl-3-[4-

(benzoyloxy)-2-hydroxy-5-[(2,3,4,6-tetra-0-benzoylhexopyranosyl)oxy]phenyl

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]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C60 H51 N5 O17

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PAGE 1-A

PAGE 1-B

— NH₂

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 126:264011 Preparation of meta-guanidine, urea, thiourea or azacyclic amino benzoic acid derivatives as integrin antagonists.

Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell; Desai,

Bipinchandra Nanubhai; Lindmark, Richard John; Rico, Joseph Gerace; Rogers, Thomas Edward; Russell, Mark Andrew; et al. (G.D. Searle & Co., USA; Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell; Desai, Bipinchandra Nanubhai; Lindmark, Richard, John). PCT Int. Appl. WO

9708145 A1 19970306, 930 pp. DESIGNATED STATES: W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (English). CODEN: PIXXD2. APPLICATION: WO 1996-US13500 19960827.

PATENT NO. KIND DATE APPLICATION NO. -----PΙ WO 9708145 A1 19970306 WO 1996-US13500 19960827 W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM CA 2230209 AA 19970306 CA 1996-2230209 19960827 AU 9671039 **A1** 19970319 AU 1996-71039 19960827 AU 702487 B2 19990225 EP 850221 Α1 19980701 EP 1996-932142 19960827 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI CN 1201454 A 19981209 CN 1996-197911 19960827 BR 9610422 BR 1996-10422 Α 19990713 19960827 JP 11510814 **T2** JP 1996-510397 19990921 19960827 NO 9800817 Α 19980424 NO 1998-817 19980226

L12 ANSWER 2 OF 17 REGISTRY COPYRIGHT 2000 ACS

PRIORITY: US 1995-3277 19950830.

RN 188811-82-7 REGISTRY

CN .beta.-Alanine, N-[3-[(aminoiminomethyl)amino]benzoyl]glycyl-3-[5-(hexopyranosyloxy)-2,4-dihydroxyphenyl]- (9CI) (CA INDEX NAME)

FS - 3D CONCORD

MF C25 H31 N5 O12

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

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1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)
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126:264011 Preparation of meta-quanidine, urea, thiourea or
REFERENCE 1:
     azacyclic amino benzoic acid derivatives as integrin antagonists.
     Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell; Desai,
     Bipinchandra Nanubhai; Lindmark, Richard John; Rico, Joseph Gerace;
     Rogers, Thomas Edward; Russell, Mark Andrew; et al. (G.D. Searle & Co.,
     USA; Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell;
     Desai, Bipinchandra Nanubhai; Lindmark, Richard, John). PCT Int. Appl.
WO
     9708145 A1 19970306, 930 pp. DESIGNATED STATES: W: AL, AM, AT, AU, AZ,
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     (English). CODEN: PIXXD2. APPLICATION: WO 1996-US13500 19960827.
     PRIORITY: US 1995-3277 19950830.
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                                                APPLICATION NO. DATE
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                                                JP 1996-510397
                                                                   19960827
     NO 9800817
                         Α
                               19980424
                                                NO 1998-817
                                                                   19980226
     ANSWER 3 OF 17 REGISTRY COPYRIGHT 2000 ACS
L12
     188811-76-9 REGISTRY
RN
CN
     .beta.-Alanine, N-[3-[(aminoiminomethyl)amino]benzoyl]glycyl-3-[2-
      (benzoyloxy)-3,5-dichlorophenyl]- (9CI) (CA INDEX NAME)
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LC
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- 1 REFERENCES IN FILE CAPILLS (1967 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 126:264011 Preparation of meta-guanidine, urea, thiourea or azacyclic amino benzoic acid derivatives as integrin antagonists.

Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell; Desai, Bipinchandra Nanubhai; Lindmark, Richard John; Rico, Joseph Gerace; Rogers, Thomas Edward; Russell, Mark Andrew; et al. (G.D. Searle & Co., USA; Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell; Desai, Bipinchandra Nanubhai; Lindmark, Richard, John). PCT Int. Appl. WO

9708145 Al 19970306, 930 pp. DESIGNATED STATES: W: AL, AM, AT, AU, AZ,

9708145 A1 19970306, 930 pp. DESIGNATED STATES: W: AL, AM, AT, AU, AZ BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (English). CODEN: PIXXD2. APPLICATION: WO 1996-US13500 19960827. PRIORITY: US 1995-3277 19950830.

PATENT NO. KIND DATE APPLICATION NO. DATE ΡI WO 9708145 A1 19970306 WO 1996-US13500 19960827 W:AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM 19970306 CA 2230209 AA CA 1996-2230209 19960827 AU 1996-71039 AU 9671039 Α1 19970319 19960827 AU 702487 B2 19990225 EP 850221 **A1** 19980701 EP 1996-932142 19960827 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI 19981209 CN 1996-197911 19960827 CN 1201454 А BR 1996-10422 BR 9610422 Α 19990713 19960827 JP 11510814 Т2 19990921 JP 1996-510397 19960827 NO 9800817 Α 19980424 NO 1998-817 19980226

ANSWER 4 OF 17 REGISTRY COPYRIGHT 2000 ACS L12 188811-50-9 REGISTRY RN .beta.-Alanine, N-[3-[(aminoiminomethyl)amino]-5-CN (trifluoromethyl)benzoyl]glycyl-3-[5-(hexopyranosyloxy)-2,4dihydroxyphenyl] - (9CI) (CA INDEX NAME) FS 3D CONCORD C26 H30 F3 N5 O12 MF CA SR CA, CAPLUS, USPATFULL LC STN Files:

- 1 REFERENCES IN FILE CA (1967 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 126:264011 Preparation of meta-guanidine, urea, thiourea or azacyclic amino benzoic acid derivatives as integrin antagonists.

Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell; Desai, Bipinchandra Nanubhai; Lindmark, Richard John; Rico, Joseph Gerace; Rogers, Thomas Edward; Russell, Mark Andrew; et al. (G.D. Searle & Co., USA; Ruminski, Peter Gerrard; Clare, Michael; Collins, Paul Waddell; Desai, Bipinchandra Nanubhai; Lindmark, Richard, John). PCT Int. Appl.

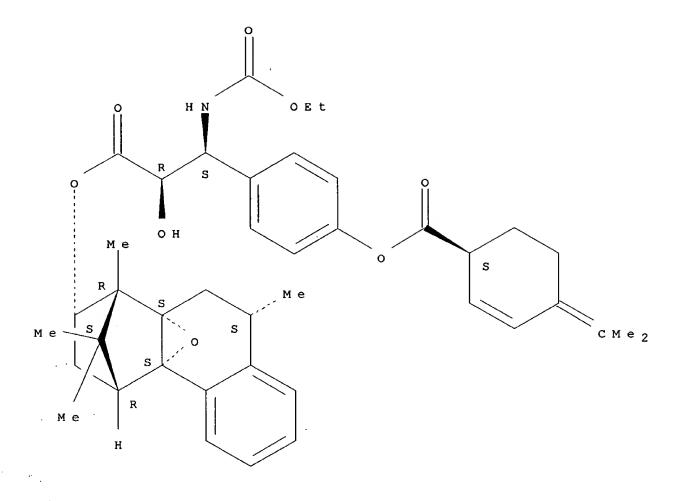
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9708145 A1 19970306, 930 pp. DESIGNATED STATES: W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (English). CODEN: PIXXD2. APPLICATION: WO 1996-US13500 19960827. PRIORITY: US 1995-3277 19950830.

PATENT NO. KIND DATE APPLICATION NO. _____ ______ 19970306 WO 1996-US13500 19960827 PΙ WO 9708145 **A**1 AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,

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                                                             19980226
L12 ANSWER 5 OF 17 REGISTRY COPYRIGHT 2000 ACS
RN
     183252-60-0 REGISTRY
CN
     Benzenepropanoic acid, .beta.-[(ethoxycarbonyl)amino]-.alpha.-hydroxy-4-
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     INDEX NAME)
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     STN Files:
                  CA, CAPLUS, USPATFULL
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Absolute stereochemistry.



REFERENCE 1: 126:7972 Preparation of borneol ester and antineoplastic and epoxyborneol ester antitumor agents. Klar, Ulrich; Graf, Hermann; Neef, Guenter; Blechert, Siegfried (Schering A.-G., Germany). Ger. Offen. DE 19513040 A1 19961002, 14 pp. (German). CODEN: GWXXBX. APPLICATION: DE 1995-19513040 19950329.

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	IE, FI								
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ES 2133953 T3 19990916 ES 1996-909153 19960329 US 5965609 A 19991012 US 1998-930009 19980313

L12 ANSWER 6 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 180263-19-8 REGISTRY

CN .beta.-Alanine, 3-[4-(cyclopentyloxy)phenyl]-N-[N-[(1,1-dimethylethoxy)carbonyl]-L-valyl]-, 1-methylethyl ester (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C27 H42 N2 O6

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 125:168644 Derivatives of beta-aminopropionic acid with a fungicidal activity. Camaggi, Giovanni; Filippini, Lucio; Gusmeroli, Marilena; Mormile, Silvia; Signorini, Ernesto; Garavaglia, Carlo (Isagro Ricerca S.r.l., Italy). Eur. Pat. Appl. EP 718280 A2 19960626, 77 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU,

NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1995-115777 19951006. PRIORITY: IT 1994-MI2156 19941021.

PATENT NO. KIND DATE APPLICATION NO. DATE PΙ EP 718280 A2 19960626 EP 1995-115777 19951006 EP 718280 АЗ 19961030 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE EP 843967 19980527 EP 1998-100374 19951006 Α1 EP 843967 20000405 В1 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, SI

AT 191317 E 20000415 AT 1998-100374 19951006

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US	5856311	Α	19990105	US	1995-553782	19951023

L12 ANSWER 7 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 170565-32-9 REGISTRY

CN Benzenepropanoic acid, .beta.-[[[4-[4-(aminoiminomethyl)phenyl]-4-methyl-2,5-dioxo-1-imidazolidinyl]acetyl]amino]-3-phenoxy- (9CI) (CA INDEX

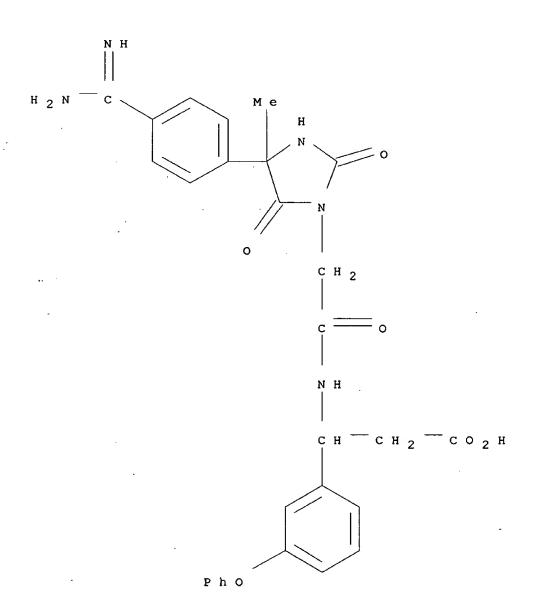
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FS 3D CONCORD

MF C28 H27 N5 O6

SR CA

LC STN Files: CA, CAPLUS



REFERENCE 1: 123:340969 Preparation of substituted 5-member-ring heterocyclic-compound blood platelet aggregation inhibitors and anticancer

agents. Zoller, Gerhard; Klingler, Otmar; Jablonka, Bernd; Just, Melitta;

Breipohl, Gerhard; Knolle, Jochen; Koenig, Wolfgang; Stilz, Hans-Ulrich (Cassella Aktiengesellschaft, Germany). PCT Int. Appl. WO 9514008 A1 19950526, 103 pp. DESIGNATED STATES: W: AU, CA, CN, CZ, FI, HU, JP, KR, PL, RU, SK, US; RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (German). CODEN: PIXXD2. APPLICATION: WO 1994-EP3491 19941024. PRIORITY: DE 1993-4338944 19931115; DE 1994-4427979 19940808. PATENT NO. KIND DATE APPLICATION NO. DATE

WO 9514008 **A**1 19950526 WO 1994-EP3491 19941024 ΡI W: AU, CA, CN, CZ, FI, HU, JP, KR, PL, RU, SK, US RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE DE 4338944 A1 19950518 DE 1993-4338944 19931115 DE 4427979 19960215 DE 1994-4427979 19940808 **A**1 AU 9479397 **A**1 19950606 AU 1994-79397 19941024 AU 693811 В2 19980709 EP 1994-930215 EP 729460 A1 19960904 19941024 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE JP 09505062 Т2 19970520 JP 1994-514176 19941024 FI 1996-2043 FI 9602043 Α 19960514 19960514 .US 5981492 Α 19991109 US 1996-640895 19960719

L12 ANSWER 8 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 147525-56-2 REGISTRY

CN Benzenepropanoic acid, .beta.-[[5-[4-(aminoiminomethyl)phenyl]-1-oxopentyl]amino]-4-phenoxy-, ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C29 H33 N3 O4

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

REFERENCE 1: 118:234488 Preparation of amidine-containing peptide mimetic compounds useful as platelet aggregation inhibitors. Garland, Robert Bruce; Miyano, Masateru; Zablocki, Jeffery Alan; Bovy, Philippe Roger; McMackins, Dudley Elgin; Rico, Joseph Gerace; Tjoeng, Foe Siong; Toth, Mihaly Vadkerti (Searle, G. D., and Co., USA; Monsanto Co.). Eur. Pat. Appl. EP 513810 A1 19921119, 66 pp. DESIGNATED STATES: R: PT. (English). CODEN: EPXXDW. APPLICATION: EP 1992-108214 19920515. PRIORITY: US 1991-702667 19910517; US 1991-792542 19911112.

PATENT NO. KIND DATE APPLICATION NO. DATE

PI EP 513810 A1 19921119 EP 1992-108214 19920515

R: PT

A 19930615 US 1991-792542 19911112 A1 19921126 WO 1992-US3573 19920506

WO 9220705 A1 19921126 WO 1992-US3573 19920506
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RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FR, GA, GB, GN, GR, IT, LU, MC, ML, MR, NL, SE, SN, TD, TG

AU 9220251 A1 19921230 AU 1992-20251 19920506

L12 ANSWER 9 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 147525-23-3 REGISTRY

US 5220050

CN Benzenepropanoic acid, .beta.-[[5-[4-(aminoiminomethyl)phenyl]-1-oxopentyl]amino]-3-phenoxy-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)

MF C27 H29 N3 O4 . C2 H F3 O2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

CM 1

CRN 147525-22-2 CMF C27 H29 N3 O4

CM 2

CRN 76-05-1 CMF C2 H F3 O2

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F — с — со<sub>2</sub> н
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REFERENCE 1: 118:234488 Preparation of amidine-containing peptide mimetic compounds useful as platelet aggregation inhibitors. Garland, Robert Bruce; Miyano, Masateru; Zablocki, Jeffery Alan; Bovy, Philippe Roger; McMackins, Dudley Elgin; Rico, Joseph Gerace; Tjoeng, Foe Siong; Toth, Mihaly Vadkerti (Searle, G. D., and Co., USA; Monsanto Co.). Eur. Pat. Appl. EP 513810 A1 19921119, 66 pp. DESIGNATED STATES: R: PT. (English). CODEN: EPXXDW. APPLICATION: EP 1992-108214 19920515. PRIORITY: US 1991-702667 19910517; US 1991-792542 19911112. PATENT NO. KIND DATE APPLICATION NO. DATE _____ EP 513810 A1 19921119 EP 1992-108214 19920515 ΡI R: PT US 1991-792542 19911112 WO 1992-US3573 19920506 US 5220050 19930615 A1 19921126 WO 9220705 W: AT, AU, BB, BG, BR, CA, CH, CS, DE, DK, ES, FI, GB, HU, JP, KP,  $\mbox{KR, $L$K, $L$U, $M$G, $M$N, $M$W, $N$L, $NO, $PL, $RO, $RU, $SD, $SE, $US$} \label{eq:kr}$ RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FR, GA, GB, GN, GR, IT, LU, MC, ML, MR, NL, SE, SN, TD, TG AU 9220251 A1 19921230 AU 1992-20251 19920506 L12 ANSWER 10 OF 17 REGISTRY COPYRIGHT 2000 ACS RN 147525-22-2 REGISTRY CN Benzenepropanoic acid, .beta.-[[5-[4-(aminoiminomethyl)phenyl]-1-• oxopentyl]amino]-3-phenoxy- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C27 H29 N3 O4

CI COM

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

REFERENCE 1: 118:234488 Preparation of amidine-containing peptide mimetic compounds useful as platelet aggregation inhibitors. Garland, Robert Bruce; Miyano, Masateru; Zablocki, Jeffery Alan; Bovy, Philippe Roger; McMackins, Dudley Elgin; Rico, Joseph Gerace; Tjoeng, Foe Siong; Toth, Mihaly Vadkerti (Searle, G. D., and Co., USA; Monsanto Co.). Eur. Pat. Appl. EP 513810 Al 19921119, 66 pp. DESIGNATED STATES: R: PT. (English). CODEN: EPXXDW. APPLICATION: EP 1992-108214 19920515. PRIORITY: US 1991-702667 19910517; US 1991-792542 19911112. PATENT NO. KIND DATE APPLICATION NO. DATE

EP 513810 19921119 EP 1992-108214 PΙ A1 19920515 R: PT US 5220050 19930615 US 1991-792542 19911112 Α WO 9220705 A1 19921126 WO 1992-US3573 19920506 W: AT, AU, BB, BG, BR, CA, CH, CS, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, PL, RO, RU, SD, SE, US RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FR, GA, GB, GN, GR, IT, LU, MC, ML, MR, NL, SE, SN, TD, TG AU 9220251 A1 19921230 AU 1992-20251 19920506

L12 ANSWER 11 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 147525-19-7 REGISTRY

CN Benzenepropanoic acid, .beta.-[[5-[4-(aminoiminomethyl)phenyl]-1-oxopentyl]amino]-4-phenoxy-, mono(trifluoroacetate) (9CI) (CA INDEX

NAME)

MF C27 H29 N3 O4 . C2 H F3 O2

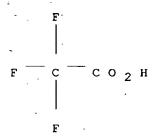
SR CA

LC STN Files: CA, CAPLUS, USPATFULL

CM 1

CRN 147525-18-6 CMF C27 H29 N3 O4

CM 2
CRN 76-05-1
CMF C2 H F3 O2



1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 118:234488 Preparation of amidine-containing peptide mimetic compounds useful as platelet aggregation inhibitors. Garland, Robert Bruce; Miyano, Masateru; Zablocki, Jeffery Alan; Bovy, Philippe Roger; McMackins, Dudley Elgin; Rico, Joseph Gerace; Tjoeng, Foe Siong; Toth, Mihaly Vadkerti (Searle, G. D., and Co., USA; Monsanto Co.). Eur. Pat. Appl. EP 513810 Al 19921119, 66 pp. DESIGNATED STATES: R: PT. (English). CODEN: EPXXDW. APPLICATION: EP 1992-108214 19920515. PRIORITY: US 1991-702667 19910517; US 1991-792542 19911112.

PATENT NO. KIND DATE APPLICATION NO. DATE EP 513810 19921119 ΡI A1 EP 1992-108214 19920515 R: PT US 5220050 19930615 US 1991-792542 Α 19911112 WO 9220705 WO 1992-US3573 19920506 A1 19921126 W: AT, AU, BB, BG, BR, CA, CH, CS, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, PL, RO, RU, SD, SE, US
RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, DE, DK, ES, FR, GA, GB, GN, GR, IT, LU, MC, ML, MR, NL, SE, SN, TD, TG AU 1992-20251 AU 9220251 **A**1 19921230 19920506

L12 ANSWER 12 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 147525-18-6 REGISTRY

CN Benzenepropanoic acid, .beta.-[[5-[4-(aminoiminomethyl)phenyl]-1-oxopentyl]amino]-4-phenoxy- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C27 H29 N3 O4

CI COM

SR CA

L12 ANSWER 13 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 110375-70-7 REGISTRY

CN Hydrocinnamic acid, .beta.-acetamido-4-(p-methoxyphenoxy)-3,5-dinitro-, ethyl ester (6CI) (CA INDEX NAME)

JUVOROV

FS 3D CONCORD

MF C20 H21 N3 O9

SR CAOLD

LC STN Files: BEILSTEIN*, CAOLD

(*File contains numerically searchable property data)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

-L12 ANSWER 14 OF 17- REGISTRY COPYRIGHT 2000 ACS

RN 102594-51-4 REGISTRY

CN Hydrocinnamic acid, .beta.-benzamido-p-hydroxy-, benzoate (6CI) (CA INDEX

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NAME)
FS 3D CONCORD
MF C23 H19 N O5
SR CAOLD
LC STN Files: BEILSTEIN*, CAOLD
(*File contains numerically searchable property data)

O
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RN-102012-12-4 REGISTRY
CN Hydrocinnamic acid, .beta.-acetamido-3,5-diiodo-4-(p-methoxypheno ethyl ester (6CI) (CA INDEX NAME)
FS 3D CONCORD
MF C20 H21 I2 N 05
SR CAOLD
LC STN Files: BEILSTEIN*, CAOLD

(*File contains numerically searchable property data)

#### 1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L12 ANSWER 16 OF 17 REGISTRY COPYRIGHT 2000 ACS RN 84878-01-3 REGISTRY

CN Benzenepropanoic acid, .beta.-[((1,1-dimethylethoxy)carbonyl]amino]-4-methoxy-3-[4-(3-methoxy-3-oxo-1-propenyl)phenoxy]-, methyl ester, (2)-(9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Benzenepropanoic acid, .beta.-[[(1,1-dimethylethoxy)carbonyl]amino]-4methoxy-3-[4-(3-methoxy-3-oxo-1-propenyl)phenoxy]-, methyl ester,
(2)-(.+-.)-

FS STEREOSEARCH

MF C26 H31 N O8

LC STN Files: BEILSTEIN*, CA, CAPLUS

(*File contains numerically searchable property data)

Double bond geometry as shown.

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 99:38691 Total synthesis of (.+-.)-chaenorhine. Wasserman,
Harry H.; Robinson, Ralph P.; Carter, Charles G. (Dep. Chem., Yale Univ.,
New Haven, CT, 06511, USA). J. Am. Chem. Soc., 105(6), 1697-8 (English)
1983. CODEN: JACSAT. ISSN: 0002-7863.

L12 ANSWER 17 OF 17 REGISTRY COPYRIGHT 2000 ACS

RN 2802-52-0 REGISTRY

CN Hydrocinnamic acid, .beta.-benzamido-3-fluoro-4-hydroxy-, benzoate (ester)

(8CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Hydrocinnamic acid, .beta.-benzamido-3-fluoro-4-hydroxy-, benzoate (6CI)

FS 3D CONCORD

MF C23 H18 F N O5

LC STN Files: BEILSTEIN*, CAOLD

(*File contains numerically searchable property data)

Novelle'

- 1 REFERENCES IN FILE CA (1967 TO DATE)
  1 REFERENCES IN FILE CAPLUS (1967 TO DATE)
- REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 Al 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

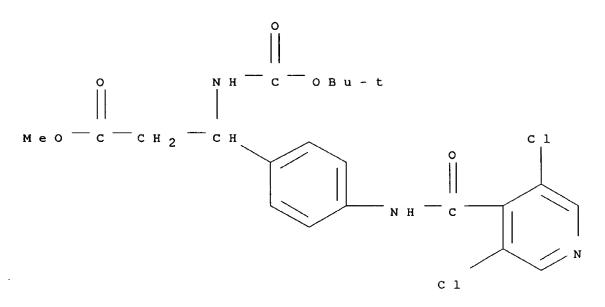
APPLICATION NO. PATENT NO. KIND DATE WO 2000032575 20000608 WO 1999-GB3986 A1 19991129 ΡI AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

- L14 ANSWER 2 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-26-6 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[(1,1-dimethylethoxy)carbonyl]amino]-, methyl ester (9CI)
  (CA INDEX NAME)
- FS 3D CONCORD

MF C21 H23 C12 N3 O5

SR CA

LC STN Files: CA, CAPLUS



1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129.

PRIORITY: GB 1998-26174 19981130. PATENT NO. APPLICATION NO. DATE KIND DATE ____ _____. PΙ WO 2000032575 A1 20000608 WO 1999-GB3986 19991129 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

- L14 ANSWER 3 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-20-0 REGISTRY
- CN Benzenepropanoic acid, .beta.-[[[(3,4-dichlorophenyl)amino]carbonyl]amino]-4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX

NAME)

FS 3D CONCORD

MF C22 H16 C14 N4 O4

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129.

PRIORITY: GB 1998-26174 19981130.
PATENT NO. KIND DATE

PATENT NO. KIND DATE APPLICATION NO. DATE

"" WO 2000032575 A1 20000608 WO 1999-GB3986 19991129

"" AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 4 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273920-16-4 REGISTRY

CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[(3-pyridinylacetyl)amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C22 H18 C12 N4 O4

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

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PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 2000032575 Al 20000608 WO 1999-GB3986 19991129

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

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- L14 ANSWER 5 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-14-2 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[(1-oxopropyl)amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C18 H17 C12 N3 O4
- SR CA
- LC STN Files: CA, CAPLUS

- 1 REFERENCES IN FILE CA (1967 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129.

PATENT NO. KIND DATE APPLICATION NO. WO 2000032575 ΡI A1 20000608 WO 1999-GB3986 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 6 OF 38 REGISTRY COPYRIGHT 2000 ACS

PRIORITY: GB 1998-26174 19981130.

- RN 273920-12-0 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[(1-phenylcyclopropyl)carbonyl]amino]- (9CI) (CA INDEX NAME)
- MF C25 H21 C12 N3 O4
- SR CA
- LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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L14 ANSWER 1 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273921-26-9 REGISTRY

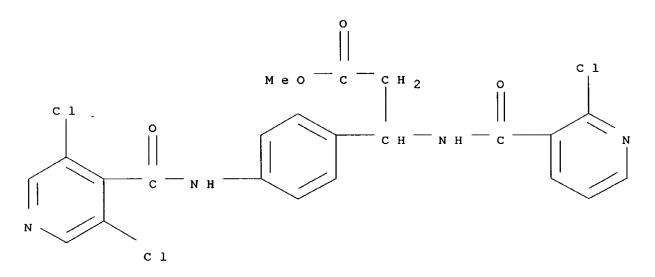
CN Benzenepropanoic acid, .beta.-[[[1-(4-chlorophenyl)cyclopropyl]carbon yl]amino]-4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

MF C25 H20 G13 N3 O4

SR CA

LC STN Files: CA, CAPLUS

SR CA STN Files: CA, CAPLUS LC



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 Al 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129.

APPLICATION NO

DATE

PRIORITY: GB 1998-26174 19981130. KIND DATE PATENT NO

	PATENT NO.		LT:	MD.	DATE			APPLICATION NO.					DATE				
ΡI	I WO 2000032575			A1 20000608			WO 1999-GB3986				6	19991129					
	₩:	ΑE,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CR,	CU,
		CZ,	DE,	DK,	DM,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,	ΗU,	ID,	IL,
		IN,	IS,	JP,	KΕ,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,
		MD,	MG,	MK,	MN,	MW,	MX,	NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,
		SK,	SL,	ТJ,	TM,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZW,	AM,
		ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM								
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		CG,	CI,	CM,	GΑ,	GN,	GW,	ML,	MR,	NE,	SN,	TD,	ТG				

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=> s 102594-51-4

## REG1stRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

KIND DATE APPLICATION NO. PATENT NO. DATE WO-2000032575 A1 20000608 WO 1999-GB3986 19991129 PΙ AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

- L14 ANSWER 7 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-10-8 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[(1-ethyl-3-methyl-1H-pyrazol-5-yl)carbonyl]amino]- (9CI)
  (CA INDEX NAME)
- FS 3D CONCORD
- MF C22 H21 C12 N5 O4
- SR CA
- LC STN Files: CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

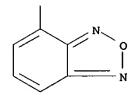
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PI WO 2000032575 A1 20000608 WO 1999-GB3986 19991129

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

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- L14 ANSWER 8 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-09-5 REGISTRY
- CN Benzenepropanoic acid, .beta.-[(2,1,3-benzoxadiazol-4-ylcarbonyl)amino]-4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C22 H15 C12 N5 O5
- SR CA
- LC STN Files: CA, CAPLUS



548/269.1

- 1 REFERENCES IN FILE CA (1967 TO DATE)
  1 REFERENCES IN FILE CAPLUS (1967 TO DATE)
- REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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PATENT NO. KIND DATE APPLICATION NO. DATE _____ _____ -----WO 2000032575 A1 20000608 WO 1999-GB3986 19991129 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM,

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- L14 ANSWER 9 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-08-4 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[(4-methyl-1,2,3-thiadiazol-5-yl)carbonyl]amino]- (9CI) (CA
  INDEX NAME)
- FS 3D CONCORD
- MF C19 H15 C12 N5 O4 S
- SR CA
- LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 Al 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

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PATENT NO.
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- L14 ANSWER 10 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-07-3 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-.beta.-[[(1-methyl-5-nitro-1H-pyrazol-4-yl)carbonyl]amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C20 H16 C12 N6 O6
- SR CA
- LC STN Files: CA, CAPLUS

### 1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 Al 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,

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PATENT NO.
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        AZ, BY, KG, KZ, MD, RU, TJ, TM
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- L14ANSWER 11 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-06-2 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-.beta.-[[(5-pyrazinyl-2-thiazolyl)carbonyl]amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C23 H16 C12 N6 O4 S
- SR
- LC STN Files: CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John: Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE WO 2000032575 20000608 WO 1999-GB3986 19991129 PΙ A1 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM; HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 12 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273920-05-1 REGISTRY

CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[(3,3-dimethyl-1-oxobutyl)amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C21 H23 C12 N3 O4

SR CA

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH,

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PATENT NO. KIND DATE APPLICATION NO. DATE

WO 2000032575 Al 20000608 WO 1999-GB3986 19991129

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L14 ANSWER 13 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273920-04-0 REGISTRY

CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[(4-methylcyclohexyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C23 H25 C12 N3 O4

SR CA LC STN Files: CA, CAPLUS

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$$C 1$$

$$C 1$$

$$C 1$$

$$C 1$$

$$C 1$$

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

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PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE WO 2000032575 20000608 WO 1999-GB3986 19991129 PΙ A1 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 14 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273920-03-9 REGISTRY

CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[(3-methylcyclohexyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C23 H25 C12 N3 O4

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

APPLICATION NO. PATENT NO. KIND DATE 20000608 WO 2000032575 WO 1999-GB3986 19991129 PI A1 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 15 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273920-02-8 REGISTRY

CN Benzenepropanoic acid, .beta.-[(cyclopropylcarbonyl)amino]-4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C19 H17 C12 N3 O4

SR CA

LC STN Files: CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp.

DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION

APPLICATION NO. DATE

PI WO 2000032575 A1 20000608 WO 1999-GB3986 19991129

AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,

MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM,

AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

- L14 ANSWER 16 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273920-01-7 REGISTRY
- CN Benzenepropanoic acid, .beta.-[(cyclobutylcarbonyl)amino]-4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C20 H19 C12 N3 O4
- SR CA
- LC STN Files: CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 Al 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

APPLICATION NO. DATE

PATENT NO. KIND DATE 20000608 PΙ WO 2000032575 **A**1 WO 1999-GB3986 19991129 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

- L14 ANSWER 17 OF 38 REGISTRY COPYRIGHT 2000 ACS
- 273920-00-6 REGISTRY RN
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-.beta.-[[(4,6-dimethyl-2-oxo-2H-pyran-5-yl)carbonyl]amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C23 H19 C12 N3 O6
- SR
- LC STN Files: CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129.

PRIORITY: GB 1998-26174 19981130.

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PATENT NO.
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                                          APPLICATION NO.
                                                           DATE
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                                          WO 1999-GB3986
    WO 2000032575
                           20000608
                                                           19991129
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        W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
            CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
             IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
            MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
            SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM,
            AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
            DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
            CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
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- L14 ANSWER 18 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273919-99-6 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[(4-piperidinylcarbonyl)amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C21 H22 C12 N4 O4
- SR CA
- LC STN Files: CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

KIND DATE PATENT NO. APPLICATION NO. DATE PΙ WO 2000032575 A120000608 WO 1999-GB3986 19991129 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

- L14 ANSWER 19 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273919-98-5 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-.beta.-[(1-oxo-4-phenylbutyl)amino]- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C25 H23 C12 N3 O4
- SR CA
- LC STN Files: CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE

""" APPLICATION NO. DATE

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- L14 ANSWER 20 OF 38 REGISTRY COPYRIGHT 2000 ACS
- RN 273919-97-4 REGISTRY
- CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[(2E)-1-oxo-3-phenyl-2-propenyl]amino]- (9CI) (CA INDEX
  NAME)
- FS STEREOSEARCH
- MF C24 H19 C12 N3 O4
- SR CA

LC STN Files: CA, CAPLUS

Double bond geometry as shown.

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroarcyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE WO 2000032575 WO 1999-GB3986 .PI A1 20000608 19991129 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 21 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-96-3 REGISTRY

CN Benzenepropanoic acid, .beta.-[[[1-(4-chlorophenyl)cyclopentyl]carbon yl]amino]-4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

MF C27 H24 C13 N3 O4

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE 20000608 WO 2000032575 PΙ A1 WO 1999-GB3986 19991129 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 22 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-95-2 REGISTRY

CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[[[2-[(2,5-dimethoxyphenyl)thio]-3-pyridinyl]carbonyl]amino](9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C29 H24 C12 N4 O6 S

SR CA

C 1

LC STN Files: CA, CAPLUS

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C 1

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

но2с —

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CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

	PATENT NO.				KIND DATE				APPLICATION NO.						DATE			
ΡI	WO 2000032575.				A1 20000608				WO 1999-GB3986						19991129			
		W:	ÆΕ,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CR,	CU,
			CZ,	DE,	DK,	DM,	· EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,	ΗU,	ID,	IL,
			IN,	IS,	JΡ,	ΚE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,
			MD,	MG,	MK,	MN,	MW,	MX,	NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,
			SK,	SL,	ТJ,	TM,	TR,	TT,	TZ,	UA,	ŬĠ,	US,	UZ,	VN,	ΥU,	ZA,	ZW,	AM,
			ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM								
		RW:	GH,	GM,	KE,	LS,	MW,	SD,	SL,	SZ,	TZ,	UG,	ZW,	AT,	BE,	CH,	CY,	DE,

DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

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RN 273919-93-0 REGISTRY

CN Benzenepropanoic acid, 4-[(2,6-dichlorobenzoyl)amino]-.beta.-[[[2-[(2,5-dimethoxyphenyl)thio]-3-pyridinyl]carbonyl]amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C30 H25 C12 N3 O6 S

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as ..., alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 2000032575 A1 20000608 WO 1999-GB3986 19991129

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L14 ANSWER 24 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-92-9 REGISTRY

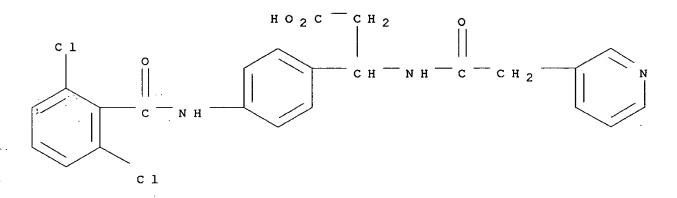
CN Benzenepropanoic acid, 4-[(2,6-dichlorobenzoyl)amino]-.beta.-[(3-pyridinylacetyl)amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C23 H19 C12 N3 O4

SR CA

LC STN Files: CA, CAPLUS



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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PATENT NO. KIND DATE APPLICATION NO. DATE 20000608 19991129 WO 2000032575 WO 1999-GB3986 PΙ A1 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 25 OF 38 REGISTRY COPYRIGHT 2000 ACS

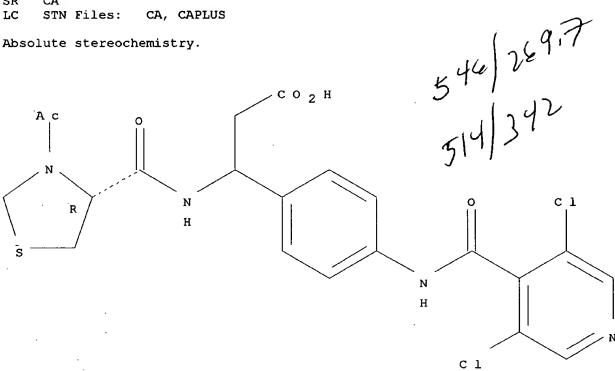
RN 273919-90-7 REGISTRY

Benzenepropanoic acid, .beta.-[[[(4R)-3-acetyl-4-CN thiazolidinyl]carbonyl]amino]-4-[[(3,5-dichloro-4pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

C21 H20 Cl2 N4 O5 S MF

SR CA



1 REFERENCES IN FILE CA (1967 TO DATE) 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. 19991129 20000608 WO 2000032575 A1 WO 1999-GB3986 PΙ AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,

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L14 ANSWER 26 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-89-4 REGISTRY

CN Benzenepropanoic acid, .beta.-[[[(4S)-3-acetyl-4-thiazolidinyl]carbonyl]amino]-4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H20 Cl2 N4 O5 S

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

- 1 REFERENCES IN FILE CA (1967 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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PATENT NO. KIND DATE

APPLICATION NO. DATE

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WO 2000032575
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    ANSWER 27 OF 38 REGISTRY COPYRIGHT 2000 ACS
L14
RN
     273919-88-3 REGISTRY
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     .beta.-Alanine, 1-[(3,5-dichlorophenyl)sulfonyl]-L-prolyl-3-[4-[[(3,5-
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     INDEX NAME)
FS
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     C26 H22 C14 N4 O7 S
MF
SR
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Absolute stereochemistry.

STN Files:

LC

CA, CAPLUS

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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PATENT NO. KIND DATE APPLICATION NO. DATE
PI WO 2000032575 A1 20000608 WO 1999-GB3986 19991129

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L14 ANSWER 28 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-87-2 REGISTRY

CN .beta.-Alanine, 1-[(3,5-dichlorophenyl)sulfonyl]-L-prolyl-3-[4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]phenyl]-2-hydroxy-, ethyl ester (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C28 H26 C14 N4 O7 S

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

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L14 ANSWER 29 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-85-0 REGISTRY

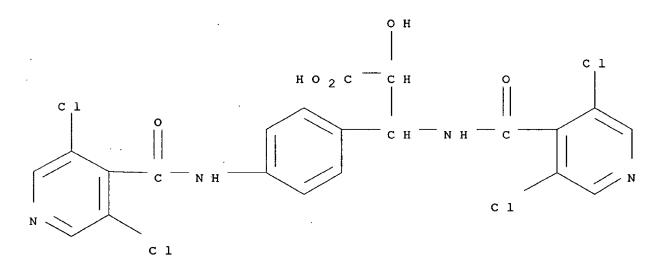
CN Benzenepropanoic acid, .beta.,4-bis[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-.alpha.-hydroxy- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C21 H14 C14 N4 O5

SR CA

LC STN Files: CA, CAPLUS



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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PRIORITY: GB 1998-26174 19981130.

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RN 273919-84-9 REGISTRY

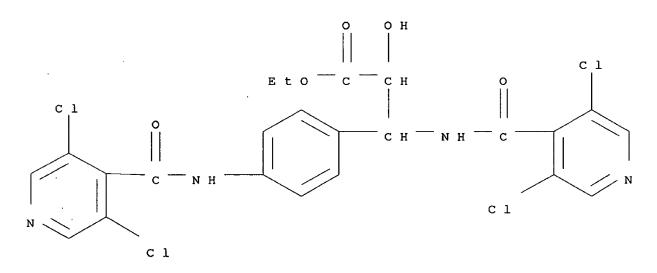
CN Benzenepropanoic acid, .beta.,4-bis[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-.alpha.-hydroxy-, ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C23 H18 C14 N4 O5

SR CA

LC STN Files: CA, CAPLUS



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

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CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129.

PRIORITY: GB 1998-26174 19981130.

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L14 ANSWER 31 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-83-8 REGISTRY

CN .beta.-Alanine, 1-[(2,6-dichlorophenyl)sulfonyl]-L-prolyl-3-[4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C26 H22 C14 N4 O6 S

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford;

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     (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129.
    PRIORITY: GB 1998-26174 19981130.
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    ANSWER 32 OF 38 REGISTRY COPYRIGHT 2000 ACS
L14
RN
    273919-82-7 REGISTRY
CN
     .beta.-Alanine, 1-[(2,6-dichlorophenyl)sulfonyl]-L-prolyl-3-[4-[[(3,5-
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     INDEX NAME)
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LC
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Absolute stereochemistry.

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. 20000608 PΙ WO 2000032575 A1 WO 1999-GB3986 19991129 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,

DATE

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L14 ANSWER 33 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-74-7 REGISTRY

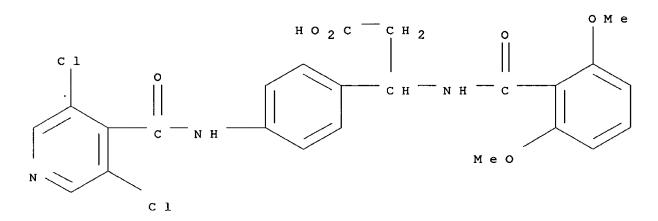
CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[(2,6-dimethoxybenzoyl)amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C24 H21 C12 N3 O6

SR CA

LC STN Files: CA, CAPLUS



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE 20000608 WO 2000032575 WO 1999-GB3986 PΙ **A**1 19991129 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,

L14 ANSWER 34 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-73-6 REGISTRY

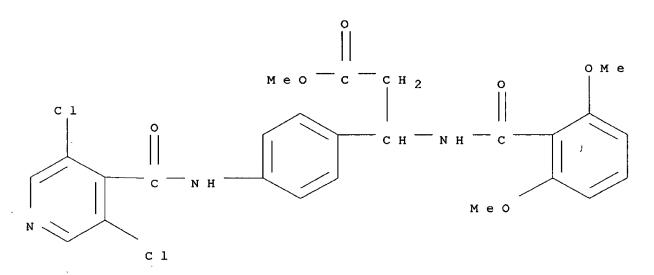
CN Benzenepropanoic acid, 4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino].beta.-[(2,6-dimethoxybenzoyl)amino]-, methyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C25 H23 C12 N3 O6

SR CA

LC STN Files: CA, CAPLUS



- 1 REFERENCES IN FILE CA (1967 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE 20000608 WO 2000032575 PΙ **A**1 WO 1999-GB3986 19991129 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,

CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

L14 ANSWER 35 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-72-5 REGISTRY

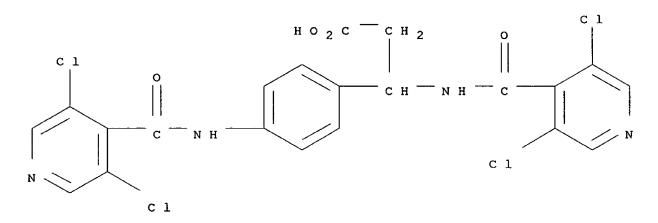
CN Benzenepropanoic acid, .beta.,4-bis{[(3,5-dichloro-4pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C21 H14 C14 N4 O4

SR CA

LC STN Files: CA, CAPLUS



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE

""" WO 2000032575 Al 20000608 WO 1999-GB3986 19991129

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

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L14 ANSWER 36 OF 38 REGISTRY COPYRIGHT 2000 ACS RN 273919-71-4 REGISTRY

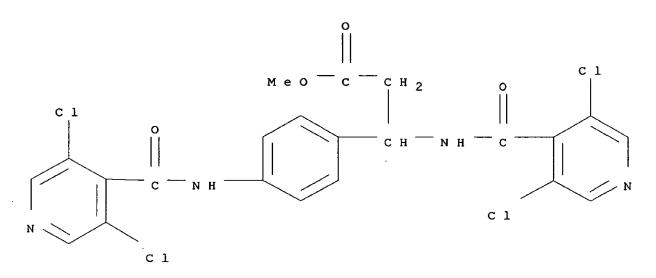
CN Benzenepropanoic acid, .beta.,4-bis[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-, methyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C22 H16 C14 N4 O4

SR CA

LC STN Files: CA, CAPLUS



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as .alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE
WO 2000032575 A1 20000608 WO 1999-GB3986 1999

WO 2000032575 A1 20000608 WO 1999-GB3986 19991129

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RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

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RN 273919-70-3 REGISTRY

PΙ

CN Benzenepropanoic acid, .beta.-[[(2-chloro-3-pyridinyl)carbonyl]amino]-

4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C21 H15 C13 N4 O4

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:30662 Preparation of N-heteroaroyl-.beta.-alanines as alpha.4 integrin inhibitors. Porter, John Robert; Head, John Clifford; Warrellow, Graham John; Archibald, Sarah Catherine (Celltech Therapeutics Limited, UK). PCT Int. Appl. WO 2000032575 A1 20000608, 66 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,

CH,

CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-GB3986 19991129. PRIORITY: GB 1998-26174 19981130.

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 2000032575 Al 20000608 WO 1999-GB3986 19991129

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,

L14 ANSWER 38 OF 38 REGISTRY COPYRIGHT 2000 ACS

RN 273919-69-0 REGISTRY

CN Benzenepropanoic acid, .beta.-[[(2-chloro-3-pyridinyl)carbonyl]amino]4-[[(3,5-dichloro-4-pyridinyl)carbonyl]amino]-, methyl ester (9CI)
(CA INDEX NAME)

CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

FS 3D CONCORD

MF C22 H17 C13 N4 O4

## REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

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L6 ANSWER 1 OF 1 CAOLD COPYRIGHT 2000 ACS

AN CA51:2748e CAOLD

TI chem. constitution and pharmacol. action in the imidazoline series

AU Novelli, Armando; Fernandez Long, P.

IT 1082-56-0 **2802-52-0** 3160-17-6 3160-18-7 17450-62-3 17450-63-4 107774-01-6 109966-27-0 110028-88-1

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FULL ESTIMATED COST 2.21 8.28

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L17 ANSWER 1 OF 2 CAOLD COPYRIGHT 2000 ACS

AN CA53:2106b CAOLD

 ${\tt TI}$  synthesis of .beta.-amino acids from aromatic hydroxy and alkoxy aldehydes

AU Rodionov, V. M.; Dudinskaya, A. A.; Avramenko, V. G.; Suvorov, N. N. IT 531-81-7 4475-29-0 5678-45-5 6049-54-3 6099-03-2 17449-03-5

68208-19-5 100397-72-6 **102594-51-4** 102872-33-3 116600-62-5 120023-66-7

L17 ANSWER 2 OF 2 CAOLD COPYRIGHT 2000 ACS

AN CA52:6260b CAOLD

TI synthesis of .beta.-diiodotyrosine

AU Rodionov, V. M.; Suvorov, N. N.; Avramenko, V. G.; Morozovskaya, L. M.

IT 830-09-1 3734-24-5 5678-45-5 6049-54-3 100884-59-1 102594-51-4 102872-34-4 102882-29-1 103855-74-9 109252-60-0

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This file contains CAS Registry Numbers for easy and accurate substance identification. Title keywords, authors, patent assignees, and patent information, e.g., patent numbers, are now searchable from 1907-1966. TIFF images of CA abstracts printed between 1907-1966 are available in the PAGE display formats.

This file supports REG1stRY for direct browsing and searching of all substance data from the REGISTRY file. Enter HELP FIRST for more information.

=> s 110375-70-7

## REG1stRY INITIATED

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L2 ANSWER 1 OF 1 CAOLD COPYRIGHT 2000 ACS

AN CA55:6431c CAOLD

TI hormones of the thyroid gland and their analogs - (V) synthesis of .beta.-thyroxine

AU Suvorov, N. N.; Dudinskaya, A. A.

IT 100379-40-6 101096-70-2 102012-12-4 102460-89-9 103386-98-7 103646-74-8 106474-14-0 **110375-70-7** 124117-92-6

=> s 102012-12-4 ··

## REG1stRY INITIATED

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L4 ANSWER 1 OF 1 CAOLD COPYRIGHT 2000 ACS

AN CA55:6431c CAOLD

TI hormones of the thyroid gland and their analogs - (V) synthesis of .beta.-thyroxine

AU Suvorov, N. N.; Dudinskaya, A. A.

IT 100379-40-6 101096-70-2 **102012-12-4**·102460-89-9 103386-98-7 103646-74-8 106474-14-0 110375-70-7 124117-92-6